ELDER CAREGIVING IN RURAL COMMUNITIES

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EXECUTIVE SUMMARY

Projected increases in older persons over the next decade, particularly in minorities and the "oldest-old" population segments (those 85+years), make elder care one of the most pressing health care issues for the 21st century. In that the characteristics of a community influence the health care of its residents, nowhere is the delivery of elder care more challenging than in rural communities, where low population density and large catchment areas combine with lack of service access and reimbursement in creating barriers to community-based elder care. Such factors contribute to institutional admission rates for rural elders that exceed those of urban communities

Community norms and cultural values as well as the strongly-held belief that families should "take care of their own problems," influence service acceptability in rural communities. Family members, friends and neighbors often fill the service gap, providing both direct and indirect care services for poor, socially isolated, underserved, frail and chronically ill elders in rural communities. Spouses, adult children, children-in-law and other extended family members are expected to provide informal care services for elders. However, out-migration of adult children to urban centers for employment often reduces the number of family members available to function as caregivers and many elders must depend upon loosely-articulated combinations of informal services from friends and neighbors. Community-dwelling and long-distance caregivers provide more than 70.1 percent of elder care services. The need to support these informal caregivers is a crucial rural health care program planning issue.

Issues of limited service access and reimbursement that create care barriers for elders also complicate delivery of assistance programs for their informal caregivers. Various surveys document the difficulties informal caregivers in rural communities experience in attempting to secure the information, disease-specific education, skill training, respite and on-going support necessary to care for a frail and impaired elder in the home. However, descriptive studies of caregiving in rural communities indicate that almost half of all caregivers do not receive assistance because they:

- Do not know they are eligible for caregiver assistance services;
- Are unaware that such programs exist in their community;
- Are too embarrassed to accept services they view as 'welfare';
- Are reluctant to seek services unless there is a "crisis"; and
- Find existing services too geographically distant to be helpful.

Demonstration projects in rural communities across a variety of states indicate that successful rural caregiver assistance programs are those: developed in collaboration with rural caregivers; endorsed by community leaders; and, staffed by persons who live in or are familiar with the cultural and social values of the community. Programs that compensate for geographic isolation through the combination of periodic home visits and innovative information technologies and

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telecommunications (e.g., peer caregiver telephone networks, telemedicine and telenursing contact systems), may offer the greatest promise for helping informal caregivers of rural elders.

Strategies for Developing Caregiver Assistance Programs in Rural Communities

Based on the diverse needs of rural caregivers and the characteristics of their social networks, successful caregiver support and training programs will:

- Offer programs suitable for both non-kin as well as kin caregivers;
- Provide a variety of informational programs for caregivers on topics such as healthy aging, symptoms and management of common chronic diseases, managing medical and drug regimens, emergency care, marital and family issues around long-term caregiving;
- Offer preventive counseling as well as supportive mental health services for distressed and depressed caregivers;
- ◆ Avoid labeling caregiver assistance programs with terms that may that make them socially unacceptable or stigmatizing for caregivers in a rural community (e.g., "Adult Day Care," "Respite Care" or Caregiver "Support" Group);
- Provide transportation services for community-based programs offerings, as well as home visitation services to individual caregivers;
- Offer caregiver "health promotion" programs in community centers e.g., blood pressure checks, pap smears, mammograms, during weekday, daytime hours so busy caregivers can combine self-care activities for themselves with a doctor's appointment for the elder during a trip into town;
- Provide telephone contact and referral service for urban-dwelling family members who are 'long-distance' caregivers for elders in the rural community;
- Make local fund-raising activities a regular part of programs; and
- ♦ Staff programs with professional, paraprofessional and volunteer personnel who are both knowledgeable about and sensitive to community culture and traditions as well as health care problems and service needs.

Implications for Helping Elders and Their Caregivers in Rural Communities

Policy makers and program makers should consider the following strategies:

- ◆ Implement needs assessment and data collection systems that accurately document the changing needs of elders and their informal caregivers in rural communities;
- ♦ Collaborate with local high schools and nearby community colleges to offer credit courses in visiting and working with frail elders and their caregivers;

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- Encourage urban health profession schools to include rural family caregiving courses in their curricula;
- Project the need for rural health care service providers by discipline and offer more post-graduation incentives for providers who practice in rural communities;
- ◆ Create articulated models of caregiver assistance programs between urban and rural health care centers:
- ◆ Develop, implement and evaluate "promising practice" models of caregiver assistance that are designed to be sensitive to the social, ethnic and geographic characteristics of rural communities;
- Expand elder care insurance coverage to include informal caregiver education and skill training; and
- Provide state and federal funding for rural health centers through "dollar matching" grant mechanisms that encourage rural communities to participate in fund raising and ownership of local caregiver assistance programs.

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ELDER CAREGIVING IN RURAL COMMUNITIES

In 2000, 35 million persons age 65 and older were counted in the United States (U.S.) Census (U.S. Census Bureau, 1990; 2000). Demographers project a dramatic increase in this elder population segment over the next decade, as baby boomers (persons born from 1946 to 1964) reach age 65. Thus, one of the most pressing issues for the U.S. in the 21st century will be care of its elders. In that informal caregivers provide more than 70% of care services for community-dwelling elders, the need to support those family members, neighbors and friends who are essential providers of the informational, emotional and instrumental aid necessary to maintain elders in their homes is a crucial community health care program planning issue. Although the proportion of elders living in rural environments varies between regions (Clifford et al., 1993), rural communities have proportionally more elders. Twenty-nine states, primarily in the Midwest and South, currently have elder populations that exceed the 12.4% national average (U.S. Census, 2000). In the South, which has a larger proportion of Blacks, almost one in three Black elders lives in a rural area (Coward & Krout, 1998). This chapter explores the barriers and challenges faced by caregivers of elders in a rural community, identifies caregiver assistance needs, highlights programs that have been successful in meeting those needs, and presents the R.U.R.A.L model for developing successful caregiver assistance programs. The paper begins by setting the socio-demographic/economic context for service provision to elders residing in rural communities.

Overview of the Rural Caregiver Population

Elders in Rural Communities

The Office of Management and Budget uses population size and density to differentiate metropolitan from non-metropolitan areas, while the U.S. Census Bureau uses settlement size and density to differentiate rural from urban settings. The terms are used interchangeably and this paper will use rural to designate these sparsely populated areas. The rural population has increased since 1950 and has been aging rapidly as a result of the desire to "age-in-place," of out-migration of youth from agricultural and mining areas, and the immigration of elders from urban areas (Siegel, 1993). Currently, 25% of elders in the U.S. live in a rural community, either alone or with their spouse.

The number of elderly persons at risk for disability, functional limitations, and chronic health problems continues to increase, creating a greater need for medical, mental health and social services, as well as economic and physical support. Indeed, an estimated 87% of the rural aged have some sort of chronic illness (Johnson, 1991), and a recent study supported by the Robert Wood Johnson Foundation concluded that at least 1 in 3 persons with chronic health conditions do not understand services they are eligible to receive, how to use these services, or who provides these services (The Institute for Health and Aging, UCSF, 1996). This is particularly true for the "oldest-old" segment of the population (those 85+years), which increased by 38% from 1980-1990, compared to only a 7% increase in the 60-84 year old cohort (Rogers, 1999). Thus, the stereotype of the "hale and hardy" rural elder is unfounded (Coburn & Bolda, 2001).

As there are fewer young and middle-aged adults in rural communities (because of employment-related migration to urban centers), fewer elders in these communities live with or

have regular access to their children. Magilvy and colleagues (2000) note that this is true even among rural Hispanic families, where cultural norms are changing because fewer adult children are available to provide care. Children and grandchildren often live far away and are unable to visit, engendering stressful life events and chronic strain for rural elders (Johnson, 1998).

The loss of young and middle-age adults also has an adverse effect on the economy of a rural community, reducing the per capita income as well as the tax dollars available for education and health care. Older adults who reside in rural communities have less formal education, are more likely to be poor, and are less likely to have health insurance coverage than their urban counterparts (Coward, McLaughlin, Duncan, and Bull, 1994; Schwenk, 1994). This is especially true for older women, the primary family caregivers in rural communities, who have the greatest economic vulnerability (Rogers, 1998). In that more rural workers are self-employed or work for small, privately-owned businesses, fewer elders in these communities will have pension, savings or investment incomes and will depend more heavily on Social Security benefits after retirement (Glasgow, 2000; Vrabec, 1995). Changes in state and federal policy will impact both rural elders and service providers, as Social Security, Supplemental Security Income and Medicare account for a major part of rural elders' incomes and provide critical support for local services (Rogers, 1999, p. 23).

Further, health care services are less accessible, less specialized and more costly to deliver in rural areas than in urban areas (Rogers, 1993). Rural elders do not have access to the same range of health care services as their urban counterparts, and fewer community-based care alternatives are available to them. And, although federal and state policies have encouraged community-based and in-home services as alternatives to institutionally-based care, rural elders continue to have a high utilization rate of nursing home services (Coward et. al., 1996).

From the statistics cited above, it would appear that the "at risk" person in a rural community is an elderly Black woman, living alone and depending on others for transportation, with only a social security pension income and whose children live at a distance. However, service providers and policy makers in rural settings must recognize that the rural aged are a heterogeneous lot, and the growing diversity that characterizes rural America defies simple definition (Buckwalter, Russell, & Hall, 1994). This argues for comprehensive and ongoing needs assessment by the Aging Network before beginning any rural caregiver projects (Lemke, Saha, Yankey, & Baenziger, 2001). (See Resources section for information on how to construct a comprehensive needs assessment survey from extant AAA databases—Dr. Jon Lemke).

Elder Care

Caregiving for an elder characteristically begins with *indirect care* services i.e., assistance with transportation, household maintenance, meal preparation and financial management. As an elder becomes more fragile and functionally limited over time, informal caregivers provide more *direct care* services such as assisting with mobility, dressing, eating and toileting. Nearby friends and neighbors may help with less personal care such as grocery shopping or transportation. However, family members: first spouses, then adult children and children-in-law and finally, other family members are expected to provide most of these services. A recent national survey of caregivers indicated 52% provided direct care services and 42% reported

giving medications and monitoring vital signs. Yet, almost two thirds of the caregivers surveyed reported they did not live with the care recipient (National Family Caregivers Association, 2000). In Barnes (1997), in her examination of the long term care needs of older rural women, notes that this segment of the population may be at particular risk, as family caregivers may not be readily available to care for them. Further, culturally bound differences in values and beliefs, especially those that underlie family patterns and intergenerational relationships, influence service efforts in rural settings (Bastida, 1988).

Kelley, Buckwalter and Maas (1999) examined and refuted three commonly held societal assumptions embedded in the social expectation of family care for elderly persons with dementia. These assumptions are that family members: 1) are not providing enough care; 2) have innate skills and knowledge about how to provide care; and 3) have access to the appropriate resources for the provision of care. Although not specific to rural caregivers, these assumptions are very relevant to this group. A national survey of caregivers of older adults found that 59% of caregivers received **no** instruction about how to perform any of the activities of care for their ill family member. In this same survey, caregivers of persons with dementia reported a great need for someone to talk with and for help understanding long term care resources (National Alliance for Caregiving, 1997). Rural elders and their families are reported to find the formal health care system too complex and confusing, especially as the number of care transitions (e.g. from hospital to home) increase (Magilvy & Congdon, 2000). The transition to care provider may be particularly difficult for adult children, especially if they must give up their role as a wage earner, resulting in financial hardship (Henderson, 1992). The complexities of providing support are compounded for long distance caregivers, who must often make decisions and implement actions without onsite assistance.

The Rural Caregiver

With the exception of affluent elders who make "amenity" moves for retirement to less-populated rural areas (Glasgow, 2000), the majority of elders who live in rural settings are commonly assumed to be long-term residents who benefit physically, emotionally and spirituality from the stability and continuity of living in a close-knit, small community. Because kin relationships and geographic proximity traditionally determine who is an informal caregiver, conventional wisdom holds that caregivers of rural elders will be their spouses, adult children, other family members and life-long, friends and neighbors, who also live in the community. This may not be the case. Exhibit 1 presents a side-by-side comparison of the commonly held assumptions of rural caregiving with the realities.

Exhibit 1: Rural Caregiving: Assumptions and Realities

Characteristic / Situation	Assumption	Reality
Caregiver-Care Recipient Kin Relationship	Rural caregivers usually are family kin (spouse, adult child, grandchildren) of the frail elder.	 Rural caregivers are likely to include a combination of: Spouses who are equally frail and impaired. Adult children who live at a distance and are "weekend" caregivers for their elderly parents. Neighbors or friends who live at some distance and have their own health problems. It may be difficult to determine who the caregiver is of the two frail spouses because of mutual levels of impairment/health conditions.
Caregiver Socio-Economic Situation	Because of longevity in the community, rural elders and their caregivers have financially stable situations, and live in comfortable, healthy situations on family farms or large, comfortable homesteads.	 The incidence of poverty among elders and their families is higher in rural when compared to urban communities. Caregivers in rural communities often give care in substandard situations lacking basic amenities (e.g., bathrooms, energy efficient heating/air conditioning systems), located in limited access areas, without public transportation.
Caregiver Support / Aid Network	Caregivers can draw on their large, extended families for additional support.	 Caregivers may be unwilling/unable to seek assistance from other family members because of the belief that caring for the elder is their "exclusive" responsibility, or they are not able to receive assistance as needed. Family conflicts around elder care issues may not be discussed with service providers because of caregiver concern that information will be shared with others in the community.
Caregiver Health / Well-Being	Because of early, healthy, physically-active lifestyles, rural elders and their caregivers have fewer physical or mental heath problems than their urban counterparts.	 Caregivers may not seek routine health screening / maintenance programs for themselves because of the belief that the elder's health problems are more important, or they are unable to ask for or find some one to care for their care recipient. Caregivers may not seek chronic illness care unless / until there is a crisis episode. Caregivers may not seek mental health services because of fear of acquiring a stigmatizing label in the community.
Caregiver Formal Service Utilization	Rural caregivers do not need and will not use formal services.	Service programs for caregivers and elders often are unavailable, fragmented, inaccessible, or subject to the vicissitudes of federal/state funding for rural initiatives. Services are not offered because they are not deemed eligible because of age or income.

The following three situations were drawn from clinic files in one southern state to illustrate caregiver needs and challenges in rural communities, including: program availability and acceptability, inadequate mental health services, and changing needs over time. Additionally, the role of caregiver attitudes, values and beliefs, stigma, and caregiver reluctance to seek help and to spend money on services, as they impede program provision in rural settings, are discussed and illustrated from the findings of a rural caregiver survey.

Situation 1: Mr. and Mrs. J.

Leroy J., a 74-year-old Black man is the primary caregiver for his wife, Ethel, who was diagnosed with Alzheimer's Disease four years ago. Mr. J. is wheelchair-bound because of a mining accident and he and his wife have been dependent on a cousin and his wife down the road for weekly grocery shopping in an adjacent community with a population of 2,500. The J's 55 year-old son and his wife drive 200-miles round-trip from the city twice a month to provide routine home maintenance chores. The Js' daughter-in-law spends her visit time doing household chores and providing minor hygiene care for her 72-year old mother-in-law (doing the laundry, filling the freezer with home-cooked meals and washing her mother-in-law's hair). The Js managed with this type of support until recently, when the 68 year-old-cousin had a stroke. The cousin's children are admitting him to a nursing home and moving his wife into the city with them. The next closest neighbor is almost five miles away. The Js' son is trying to get his parents to move into the city too. Although the elder Mr. J. is having difficulty managing his wife's increasing confusion, he is adamant that they not move from this house where they have lived for 50 years.

One of the first challenges in developing rural caregiver assistance programs is ensuring program services are available and acceptable to the caregivers. The Js exemplify caregiving situations common in many rural communities, where increasing health problems of an ancillary caregiver disrupts the relative stability of the home situation and the primary caregiver, who can be as impaired as the elder for whom they are caring, is resistant to change. In this situation, while they looked for someone in the community who could do the weekly shopping, the J's son and his wife increased their trips from the city to once a week. The situation changed three months later, when the elderly Mrs. J died from a cerebral aneurysm and Mr. J finally agreed to move into the city.

Situation 2: The G. Brothers

B.G. and R.G. are unmarried brothers who live together on the family farm. B.G. is 68 years old and R.G is 65. B.G. has been the primary caregiver for his younger brother, a chronic schizophrenic, since the death of their parents 15 years ago in an automobile accident. Their closest neighbor lives three miles away. The G. home is at the end of a long, winding gravel road, but bad weather often makes the road temporarily impassible. A married. 76 year-old sister who lives 50 miles away in another small farming community, contacted the regional community health center to ask that someone stop by the family homestead to 'check' on her brothers, who have stopped answering the phone. When a volunteer goes out to the house, B.G. answers the door but refuses to admit the visitor. He is dirty, unkempt and has bruises on his face.

In this situation, R.G. had stopped taking the psychotropic drugs necessary to control his schizophrenia and his increasingly frail elder brother became the object of his abuse. Based on the visitor's report to the community health center, welfare workers came out to the house and R.G. subsequently was admitted to the closest hospital as an acute psychiatric admission and finally to the state hospital as a psychiatric in-patient. One of the most frequently unmet needs in rural communities is for preventative and supportive mental health services (Neese, Abraham, & Buckwalter, 1999). Rural communities are less likely to have mental health professionals available in the community or to have the resources to offer in-service training on mental health techniques to their staff (Bane, 1997). Sources of mental health services in rural areas are often limited to community mental health centers, state hospitals, geographically scattered private practitioners, physicians and clergy. Utilization rates remain extraordinarily low (Neese et al., 1999).

Bane (1997) notes that service delivery problems in rural areas are also adversely affected by the passiveness of the system; that is, the system waits to be contacted rather than case-finding those individuals and their caregivers who are in need of assistance. One exception is the Gatekeeper program developed by the late Ray Raschko at the Spokane Community Mental Health Center Elderly Services. The Gatekeeper approach uses telephone information and referral, multidisciplinary in-home evaluation, treatment, and case management. An established system of rural community "gatekeepers," including meter readers, county assessors, and postal workers, helps identify residents in need of services. There is a pressing need for state-sponsored mental health training for rural health care providers (e.g., through outreach and distance learning programs) as well as for 'mobile' mental health providers who can take services out to homebound rural caregivers who are caring for an elder with a chronic mental illness. Mental health outreach programs in rural Iowa and Virginia are described later in this paper, and serve as promising practice models for overcoming the challenge of inadequate mental health services and identifying persons who are isolated and in need.

Situation 3: Mrs. S.

Mrs. S. is an 86 year-old woman who has lived alone in the family home since her 88 year-old husband's death a year ago. While Mrs. S. initially was able to drive into town, she has not driven the car in the last few months because she had a couple of minor accidents. One occurred on a country road late at night when she misjudged a turn and ended up in a ditch. She was forced to stay in the ditch all night, until a school bus driver on her way to work found her the next morning. Now Mrs. S' activities are limited to mowing her lawn and working in her garden. Mrs. S' caregiver is her 82 year-old-sister Barbara, who lives across the road. While Barbara calls to check on Mrs. S. every day, her time is limited because she must care for her own husband, who has a progressive neuromuscular disease. Barbara also fell on an icy patch last year and fractured her arm in three places resulting in physical limitations. Mrs. S' only other living relative is a daughter who is a Christian missionary in Africa. Over the past few weeks, Barbara has begun to notice that Mrs. S. rambles a great deal and has a limited attention span. She is concerned about her sister but unsure what to do.

A dominant feature of elder caregiving networks is their continuous evolution over time (Peek, Zsembik & Coward, 1997). While most elders initially require only limited assistance, after reaching a certain level of frailty the need for assistance increases dramatically. In rural communities, the type of assistance needed may not be readily available or, may require a combination of caregivers and services be 'cobbled' together to keep the elder in the home. In the case of Mrs. S., over the span of a single year, she progressed from being her elderly husband's caregiver to independently living alone with daily, minimal support phone contacts from her sister, to being a frail elder who may or may not be able to continue living in her own home

Appropriate services would be those designed with the recognition that need for different services could vary over time; care recipients and caregivers could seesaw between levels of professional care or care environments and community-based informal support systems. Transitions between levels of care should be as "seamless" and "fluid" as possible, which entails information transfer and exchange across care environments and providers (Chalifoux, Neese, Buckwalter, Litwak & Abraham, 1996, p. 477). Lemke and colleagues (2001) report that service usage varies across counties even within the same rural region, in part because of lack of consistency in definitions and eligibility requirements for similar programs. He argues that potential caregiver clients must be followed from initial screening, which should be available upon demand. Also, agencies must be able to track caregivers, household composition and functional status over time and the corresponding data must be maintained and archived (Lemke et al., 2001).

Factors Impeding Service Provision to Rural Caregivers

According to Van Hook (1987), the rural elderly are often provided with "scaled down urban service models that fail to meet their needs or are insensitive to the real differences between urban and rural areas" (p.13). To be successful, services must mesh with other local programs and informal helping networks, and service providers must understand and be sensitive to the rural value system and social ecology of the area (Buckwalter et al., 1994). Geographic distance from urban centers often precludes access to community-based assistance programs and most rural communities are limited in fiscal resources and infrastructure (e.g. program planners, trained workers, transportation) necessary to develop their own community-based programs (Lemke, et. al., 2001). This often results in barriers related to: awareness, availability, access, affordability, adequacy, appropriateness and acceptability (Krout, 1994; Williams, Ebrite & Redford, 1991). Rural residents may be unaware of services in their area or, the lack of specific service availability in the community may decrease service affordability and increase the time required to access those services. Locally available services may be fragmented or of lesser quality (adequacy) or may not have the necessary articulation with urban referral centers. Further, services may not target the health care needs of residents (appropriateness) or be acceptable to the cultural norms and ethnic beliefs of community residents. Connell et al., (1996) noted that even when transportation was available, caregivers in their focus group interviews reported that "the need to travel long distances effectively eliminated the intended benefit of support services (e.g. respite/adult day care, support groups) because the caregiver's "free time" was spent in the car" (p.23).

An investigation of the needs, resources and responses of n=107 rural caregivers of persons with Alzheimer's Disease (Buckwalter et al., 1994) revealed that only 51% of caregivers used any community-based services, with costs ranging from \$16-\$850 per month and averaging \$73. The majority of costs for services was borne by the caregivers themselves, who reported that cost was a major barrier to service utilization, and that they were "saving their money to buy care in a good nursing home." Another reported barrier was the concern that confidentiality would be comprised. Others stated they would not use any service where they accepted aid from agencies because they felt this was "too close to charity" and that "people should be self-sufficient" and "take care of their own problems" (p. 312). In Iowa's Mental Health of the Rural Elderly Outreach Project, many rural elders and their caregivers believed that they should be able to handle problems themselves and viewed mental health assistance as a sign of personal weakness, if not defeat (Smith, Buckwalter, & DeCroix-Bane, 1997). Similar attitudinal barriers were reported by Connell et al., (1996, p. 23), who noted that rural family caregivers in focus group interviews expressed reluctance to seek community-based services because they were seen as "hand-outs" or "welfare." These caregivers felt they should be selfreliant and solely responsible for the care of their older family members. These values and beliefs, and the stigma associated particularly with the use of mental health or counseling services, accounted in large part for the low use of formal services in this rural population. Other program-related barriers reported included poorly publicized programs, those perceived as having "too much red-tape," and programs with certain restrictions such as those that are means-tested. These barriers are consistent with those reported by Collins and colleagues (1991) who noted that denial of symptoms and reluctance to seek help can be attributed to feelings of shame, stigma, fear of institutionalization and suspicion of the health care and service systems. Ageism and misconceptions about illnesses, especially dementia, among rural

health care professionals also diminishes the likelihood that caregivers will receive adequate information and referral for needed services (Connell et al., 1996).

Connell et al., (1996, pp.19-22) developed a Community Outreach Education Model Program (COEP) for dementia caregivers in rural Michigan. Using focus group interviews with family caregivers and service providers, they identified a number of structural and attitudinal barriers to effective service delivery. Structural barriers affected help-seeking behavior and the experience of being a caregiver. These barriers included: 1) Lack of coordination in the service delivery system; 2) Cost of services; 3) Service agencies that are overburdened, understaffed, or unavailable; 4) Distance and transportation; 5) Reimbursement policies for services are too restrictive; 6) Lack of access to comprehensive diagnostic and assessment services; and 7) Family physicians do not always make referrals for services. Caregivers also reported frustrations related to the utilization of needed services, such as being put on a waiting list for services when they had an immediate need, and difficulty understanding complex reimbursement policies. Attitudinal barriers included: 1) Stigma and guilt about seeking help and receiving services; 2) Value of self-reliance; 3) Belief that family members should be responsible for care; 4) Reluctance to seek services until a crisis occurs; 5) Denial of symptoms; and 6) Ageism. This same team also noted a number of community strengths to build upon in designing services for rural caregivers, including: 1) Community action and cooperation; 2) Close knit ties and long-established roots; 3) Dedication and caring for residents in rural areas; and 4) Strong cultural identity.

Despite these acknowledged barriers, rural caregivers in the Buckwalter et al. survey (1994) were highly motivated to undertake a difficult and burdensome role, and many reported a deep sense of personal satisfaction and growth from the caregiving experience. Data from the Caregiver Burden Scale indicated a strong sense of moral obligation and personal desire as primary motivations for becoming a caregiver, and reflected cultural and religious values of the region. For example 92% of respondents indicated that "Caring is the Christian thing to do" and a similar percentage reported that "God helps them in their caregiving efforts." These findings are supported by more recent research conducted by Martinez (1999) and Davis and Magilvy (2000), who found that rural elders felt they had a responsibility to help themselves, and also felt, that along with God's help, they could survive life's challenges.

These attitudes may explain, in part, the reluctance of many rural caregivers to seek help. Robinson (1988; 1990) has developed a successful social skills training model for caregivers that enables them to learn specific skills they need in order to mobilize help from their social networks. The training program focuses on assertive social skills such as how to give instructions when help is offered, how to make a request for help, how to express appreciation for support, as well as how to say "no." Individual counseling sessions are augmented by written materials reinforcing key points covered during the counseling sessions. The model has been found to increase social support and lower burden for caregivers.

Issues in Rural Case Management

Case management is designed to help clients find the most appropriate and least costly services that will meet their needs. Parker and colleagues (1992) published an extensive review of the issues and challenges related to providing case management services to families living in rural

areas. They detail the most critical health care delivery and financing problems faced by rural communities, including third-party payer restrictions, reimbursement levels, provider recruitment and retention, competition with urban providers, emergency medical services, and regulations and paperwork. Additionally, differences between rural and urban case management are highlighted, focusing on lack of available and accessible services, financial constraints, staffing issues, poor discharge planning, and turf issues. These same themes are echoed throughout the rural case management literature. Krout (1993) has cogently argued that case management is a process that can play an important role in overcoming many of the service delivery barriers cited previously by containing costs, increasing awareness of and access to in-home services, targeting resources, preventing costly and unneeded institutionalization, coordinating various aging services, and, importantly for this paper, supporting family caregivers.

More recently (1997) Kraut analyzed data from a national sample of n=356 agencies regarding barriers to providing case management services to rural older persons. To obtain the sample, a list of 1,201 rural case management providers was developed through phone calls to state health and social service departments and state units on aging officials nationwide, who were read the definitions of case management and rural, and who were then asked to provide contact information for agencies meeting those definitions within their state. For purposes of this research, only agencies that identified a primary service area as rural, open country, small town, or village, or some combination of those settings were included in the final sample (Krout, 1997, p. 144). Several areas were consistently reported by respondents as problematic, including lack of resources to pay for case management and services, a lack of services, lack of transportation, and too many regulations. Demand for documentation made by various government agencies is an issue that plagues rural health care and service providers, whether it be for purposes of claims submission, quality assurance, or accreditation. Providers are generally not reimbursed for these paperwork demands, and must absorb the cost of maintaining staff to provide the necessary documentation (Parker et al., 1992, p. 48). Rural providers also complain that federal and state regulators as well as peer review organizations fail to adequately consider the applicability of various regulations to rural areas when implementing policies. Finally, many rural providers suggest that not only are the regulations imposing, but the rules change so frequently that it is difficult to keep track of them (Parker et al., 1992, p. 48).

Review of the Literature on What Supports Rural Caregivers

Caregiver Assistance Programs

Informal caregiving for a frail elder typically lasts for eight years or more (MetLife, 1999), and the toll of caregiving has been well documented. Numerous studies indicate informal caregivers experience role strain, negative mood and greater physical health declines than agematched non-caregivers, as well as more marital discord, family dysfunction and social isolation and loneliness (c.f., Aneshensel, Pearlin & Schuleer, 1993; Gwyther, 1995; Johnson, 1998; Lieberman & Fisher, 1995; Vedhara et al., 2000). These adverse outcomes are compounded by losses from the social support network, and limited knowledge of local resources (Maglivy & Congdon, 2000). Indeed, caregiver strain is a major precipitant of

premature and inappropriate institutionalization among rural elders, especially those who perceive they have no alternative care options (Congdon & Magilvy, 1998).

Regardless of whether they provide indirect or direct care, informal caregivers of elders in rural communities need a core set of skills that will enable them to accurately monitor and interpret symptoms, successfully manage medical regimens, provide hands-on care, find and utilize appropriate resources and make sound caregiving decisions, all while providing affective support and encouragement to the chronically-ill or disabled person (Schmall, 1995; Schumaker, Stewart, Archbold, Dodd & Dibble, 2000). Thus, caregiver assistance programs traditionally combine information on aging and chronic illness with skill training on providing home care, as well as affective support, counseling and periodic respite for the caregiver (c.f., Toseland, Smith & McCallion, 2001, for a discussion of the content of caregiver training programs).

The Role of Technology in Delivering Services to Rural Caregivers

Redford and Parkins (1997) provide an excellent review of the promise of communication and information technologies to expand the reach of case managers and improve the coordination, access to, and quality of care in rural communities. Due to increasing pressures on service infrastructures, case managers face constant challenges in finding, obtaining and monitoring services for rural clients, and educating caregivers around their care decisions. Redford and Parkins (1997) suggest that telecommunications and information technologies may be one answer to these access challenges and the effects of geographic isolation. They list the following benefits of telehealth technologies for **Case Managers** (p. 156):

- Reductions in travel time to adequately assess and monitor clients;
- Increases in the numbers of clients that can be effectively monitored;
- Opportunities to more frequently conduct educational sessions and provide technical and emotional support to homebound care recipients and their caregivers; and
- ♦ The ability to readily access through the Internet, client records, educational and informational materials, communications with colleagues, and other activities that decease isolation.

And benefits for **Consumers**, which, for purposes of this paper are caregivers:

- ♦ Closer contact with case managers;
- Opportunities for interaction with others in similar circumstances;
- Timely access to face-to-face contact when needed for assistance or support;
- Reductions in time and energy needed to travel for health monitoring;
- Assistance with the supervision and monitoring of in-home workers; and

♦ More opportunities to attain information and education to manage their own and another's care.

An example of a successful technology-based intervention was that developed by Brennan and colleagues (1995in which they established and evaluated a computer network (ComputerLink) to meet the educational and support needs of in-home caregivers of persons with dementia. The computer network provided information (e.g. a self-care encyclopedia), communication (e.g. an anonymous question and answer segment and a public bulletin board forum); and decision-support functions (among caregivers and a clinical expert). Outcomes included a significant increase in caregiver confidence in decision-making and social support.

Preliminary data from in-home telemedicine demonstration projects in rural Kansas (Lindberg, 1997) suggest that telemedicine can effectively provide health care for elderly and disabled persons, and assist their caregivers. As the literature indicates that rural elders are less likely than urban elders to use formal, in-home long term care services (Kenney, 1993), innovative technology-based service delivery models like the Kansas project, may be particularly important service delivery mechanisms for rural care providers. The next section of this paper briefly reviews some of the approaches and programs that have been effective, or show promise of being effective, in delivering needed services to rural caregivers.

Promising Practices

Mobile Outreach Programs

In 1978, the President's Commission on Mental Health noted, "Rural communities tend to be characterized by higher than average rates of psychiatric disorders, particularly depression, by severe intergenerational conflicts, by an exodus of individuals who might serve as effective role models for coping, by an acceptance of fatalistic attitudes and minimal subscription to the idea that change is possible" (p.1164). Regrettably, not much has changed over the past 23 years. Together with the often-fierce sense of self-determination, independence, dignity, privacy, and hardiness characteristic of many rural Americans (Lee, 1993), access issues and the pronounced stigma associated with mental illness, elders most at risk do not present themselves for traditional mental health services. Fear of being labeled as crazy, of being shunned by friends and neighbors, or being "put away" in an institution prevents many rural residents and their caregivers from receiving needed mental health services and supportive assistance. Thus psychiatric problems among rural residents often go undiagnosed and untreated (Abraham, Buckwalter, Snustad, Smullen, Thompson-Heisterman, Neese, & Smith, 1993; Neese et. al., 1999).

Interdisciplinary psychogeriatric outreach models in rural Iowa and Virginia have been shown to be effective (as well as cost-effective) in delivering services to geographically and socially isolated elders and their caregivers (Abraham et al., 1993; Buckwalter, Smith, Zevenbergen, & Russell, 1991; Smith & Buckwalter, 1999). Both programs are community partnerships involving the local Area Agency on Aging and the community mental health system. (For a detailed description of the key components and a comparison of these outreach models, see Abraham et al., 1993, p. 206. For more information on the structure of the Mental Health of the Rural Elderly Outreach Project see Smith & Buckwalter, 1999, and for data on

outcomes of the project see Buckwalter et al., 1991). These models also emphasize the need for strong coordination and cooperation among mental health, medical and social service providers; maximizing limited resources; assuring continuity of care; and using professional, paraprofessional, and lay personnel. The mismatch between the needs of caregivers of the mentally ill rural elderly and the services available to them will not disappear in the foreseeable future, raising the short-term service issue regarding improvement of accessibility to local and regional resources, and the long-term policy issue of determination of reasonable levels of resource development, allocation, and equity. The problems of transportation, poverty, lack of trained personnel, low population density, and the large catchment areas will continue to make the delivery of mental health services more difficult in rural areas. Outreach models can make substantial gains in overcoming these problems and providing needed services to rural elders and their caregivers. Individualizing these programs to reflect local culture, geography, need and resources improves both the delivery of care and the ability of programs to be sustained over the long term (Abraham et al., 1993, p 210).

An option for caregiver support in rural settings often comes from faith-based initiatives. Under the leadership of Dr. Karen Robinson, the University of Louisville School of Nursing developed a successful dementia-specific **Volunteer Caregivers Program (VCP)** that is an outgrowth of the Volunteer Interfaith Caregivers of Kentucky. The idea for this support program began when representatives from various religious congregations and a local Alzheimer's Disease chapter joined together and developed a vision to train volunteers to provide in-home respite to keep caregivers connected to their support groups. Since 1994 the VCP has provided support services to more than 100 caregivers at no charge. Program services include assessment, care planning, education, evaluation, information and referral, and volunteer support. The VCP also provides free public services such as memory screening and community education programs. (See Resources section for contact information)

Building a Seamless Delivery Dementia Care System in Rural Iowa (Dr. Janet Specht, PI, Dr. Geri Hall, Co-PI and Ann Bossen, Project Director) is a collaborative project between the Iowa Department of Elder Affairs, the University of Iowa's College of Nursing and Center on Aging, Area Agencies on Aging, Alzheimer's Association chapters and ResourceLink of Iowa. The three-year (2000-2003) Administration On Aging funded project (grant # 90AZ2366) is designed to provide expanded in-home services to rural Iowans and their caregivers who are affected by Alzheimer's Disease and related disorders (ADRD). The project uses a community organization model, developed by the Big Sioux chapter of the Alzheimer's Association, to help rural communities develop care systems that meet their unique needs in eight rural counties throughout Iowa. Two assumptions underlie this approach: 1) If services are to be accepted within rural communities, they must be designed by the community; and 2) If the community values the services, they will refer neighbors to them. Other successful outreach programs have also been based on a community development model (see The Community Outreach Education Program, as described by Connell et al., 1996), which is a process of working collaboratively with community members "to assess the collective needs and desires for healthful change and to address these priority needs through problem solving, utilization of local talent, resource development and management" (Lassister, 1992, p. 30). Lassiter (1992, pp. 30-31) sets forth five tenets of community development that are of special importance in rural areas: 1) Citizen participation and partnership are essential for community improvement and growth; 2) The focus of work will be on local concerns; 3) Citizen groups are utilized in

community development; 4) Implementation is suitable to the locality; and 5) Process outcomes for the community are as important as task undertakings.

The "Seamless Delivery System" project is also designed to demonstrate the effects of an inhome nurse care managed delivery system on care recipient and caregiver well being. Four (experimental) of the 8 counties receive nurse care managers (NCM) who are specially trained to provide in-home services, education and support to persons with ADRD and their caregivers, with a particular emphasis on building capacity in the caregiver by focusing on the well-being and stressors clients identify. The NCMs work with the local case management system to coordinate available services, refer clients to funding sources, assist with community development programs and report to the AAAs. There is also a telecommunications back up system, ResourceLink of Iowa, available for both control and experimental counties. Individualized services are offered to both persons with dementia and their caregivers, and dementia-trained nurses work with care recipients and caregivers. The four control counties have a local project facilitator (LPF) who is the referral point for services, completes intake forms, and refers clients to the existing Case Management System. Caregiver outcomes include Caregiver well being, endurance, and stress. (See Appendix for Client Intake Forms for this project, especially Part 2: Information about the caregiver). The investigators have identified a number of structural and philosophical barriers to implementation, including reticence to allocate services to caregivers assessed as having too many resources or too much income. At present, only baseline quantitative data has been analyzed, so comparisons between control and experimental counties are premature. However, several unique observations have emerged that may prove useful in future programming efforts. Project implementers and evaluators report that:

- ♦ When the NCM is from the area she/he serves, the project is more successful. In many cases, the nurse has become a bit of a local celebrity, achieving the status one aging network professional described as "a trusted relative," and has been easily integrated into the community's perception as a "helping professional." This integration has fostered referrals from sources such as the local bank, churches, service and social clubs, and people in the rural town.
- ♦ Several of the counties have enthusiastically embraced the community action programming. The fact that it was developed within and by their community exclusively has become a real source of pride. Programs that are clearly identified with community needs are more aggressively pursued than those that are developed by "outsiders" and "imposed" on consumers in the rural communities.
- ♦ An unanticipated problem that has emerged in this project has to do with devaluation of the younger working caregiver. Some members of the community and the aging network have expressed concerns over providing services to someone who "works and has a salary," feeling younger, working caregivers aren't "justified" in receiving help. Clearly more education aimed at changing these attitudes is required—e.g., education illuminating the adverse psychosocial and financial consequences that may ensue should the caregiver be forced to quit employment in order to be eligible for services. One approach that was effective was to emphasize to members of the aging network that should the working caregiver have to resign, he/she would no longer be

contributing to the county tax base and might also have to forfeit retirement earnings; however, attitudes denying services to employed caregivers in rural areas persist.

In-Home Caregiver Support Programs

A number of in-home caregiver support programs have demonstrated an array of positive outcomes for caregivers. Buckwalter and colleagues (NINR, 1992) conducted a 4 year multisite rural caregiver study to test the effectiveness of an in-home caregiver training intervention based on the **Progressively Lowered Stress Threshold (PLST) Model**. The intervention provided 5 hours of training to facilitate caregivers' knowledge of dementia and development of competence in problematic behavior management. Caregivers in both the experimental and control groups also received referrals for support groups and case management services. Findings revealed that caregivers who received the in-home training intervention felt better about their situation, had more satisfaction with the caregiving experience, an increased sense of mastery, as well as decreased levels of burden, uncertainty and depression. They were also less bothered or upset by behaviors of the care recipient. Of note is that the "Seamless Delivery System" project discussed above uses elements of the PLST model as part of the in-home service provided by NCMs.

Archbold and associates (1995) tested the **Preparedness, Enrichment, Predictability (PREP)** system for in-home family caregivers of elders. Through a Medicare waiver provision, families in the experimental PREP group received 3-6 months of care from PREP nurses, including systematic assessment, family focus, local and cosmopolitan knowledge, individualized interventions using multiple strategies, therapeutic relationships and transitions. Subjects in the control group received standard HMO and in-home health agency care. Caregivers who received the PREP system reported higher care effectiveness and overall usefulness of the in-home staff. Hospital costs associated with the PREP group were also lower (\$2775 vs. \$6929).

National Family Caregiver Support Program

As a result of the National Family Caregiver Support Program (NFCSP) initiative, a number of programs targeting both middle age rural caregivers in the workforce as well as older spouses are in development, or currently underway. The following are two examples of programs underway in Iowa. Under the leadership of the Iowa Association of Area Agencies on Aging (with Betty Grandquist, Coordinator), a new, multi-faceted project is underway to assist caregivers in rural settings. This project is a collaborative effort between the Iowa Association of AAAs, the State Unit on Aging, and the AAAs. It combines a case management approach with screening and referral to the nearest AAA where a "Family Caregiver Expert" takes over. This expert has a background in human services and aging issues, and is knowledgeable about resources in the communities served. A software program (ESP) developed by the Atlanta Regional Commission has been purchased which will provide a standard information and assistance network, a directory of resources, and match caregivers' needs to available options. Data will be collected statewide at the Association office, although each AAA will develop support services responsive to the needs of caregivers in their area. The Family Caregiver Expert is expected to take special classes on how to effectively use this software. In addition, a toll-free number has been established in the movement toward a single point of entry for

services. An Iowa Family Caregiver Project Web page has been developed and a marketing and education plan is being established to provide consistency across the network.

Another "Promising Practice" to be highlighted is The Family Caregiver Support Program, funded by the Heritage Area Agency and operated by Elder Services Incorporated, Iowa City, Iowa. It serves family caregivers in a seven county region and is comprised of two primary components: Information and Assistance (I & A), and Family Caregiver Counseling. The I & A Specialist receives inquiries from a nationwide toll free telephone number and provides callers with information about appropriate state or local caregiver resources. The Family Caregiver Counseling Specialist works one-on-one with family caregivers in their home to assess their needs, develop a plan to address problems, and enable caregivers to successfully maintain their role. Typically, a short-term counseling model (usually 1-6 meetings) supports caregivers during times of transition and assists them in making important decisions, although crisis intervention services are also available. The Counseling Specialist may provide assistance in the form of I & A, skill building (e.g. communication, caregiving skills), short term counseling, family mediation, and arrangement of appropriate community services and resources. Information is targeted to each caregiver's unique needs, and help is provided to alleviate their sense of isolation and to feel supported in their role. Family Caregiver Support services are funded by the NFCSP and are provided free of charge. The majority of referrals come from case managers who oversee services to care recipients involved in the Case Management Program for the Frail Elderly. The Counseling Specialist addresses the caregiver's needs, which may go unnoticed in the traditional case management program. Other referral sources include health care centers, aging service providers and members of the community at large.

Practice Implications for the Aging Network

The R.U.R.A.L Model

Based on the diverse needs of rural caregivers and the characteristics of their social networks, successful caregiver support, and training programs will:

- ◆ Offer programs suitable for both non-kin as well as kin caregivers (e.g., "How to start a Neighborhood Friendly Visitor Program");
- Provide a variety of informational programs for caregivers on topics such as healthy aging, symptoms and management of common chronic diseases, managing medical and drug regimens, emergency care, marital and family issues around long-term caregiving;
- Offer preventive counseling as well as supportive counseling services for distressed and depressed caregivers;
- Offer community programs that avoid labels that may make them socially unacceptable or stigmatizing for caregivers in a rural community (e.g., "Adult Day Care," "Respite Care" or Caregiver "Support" Group);

- ◆ Provide access to transportation services to community-based programs offerings, as well as home visitation services for individual caregivers;
- Offer caregiver "health promotion" programs e.g., blood pressure checks, pap smears, mammograms, during weekday, daytime hours so busy caregivers can combine self-care activities for themselves with a doctor's appointment for the elder during a trip into town;
- ◆ Provide telephone contact and referral service for urban-dwelling family members who are "long-distance" caregivers for elders in the rural community;
- ♦ Make annual fund-raising activities a regular part of program so as to insure local investment/ownership in the project, which is critical to its success; and
- Staff programs with professional, paraprofessional and volunteer personnel who are both knowledgeable about and sensitive to community culture and traditions as well as health care problems and service needs.

Based on the descriptive studies and projects highlighted in this paper, Exhibit 2 lists program characteristics and persons to involve for planning successful rural caregiver assistance programs.

Exhibit 2: R.U.R.A.L Caregiver Program Planning Model

	Desired Program Characteristic	Program planners must ensure that rural caregiver assistance programs:
R	<u>R</u> elevance	Involve caregivers in identifying program service needs and program relevance
U	<u>U</u> nity	Integrate new program offerings with existing community services to insure the new program does not compete with or duplicate existing programs
R	<u>R</u> esponsiveness	Are responsive to the ethnic and cultural identity and traditions of elders/caregivers/ residents in the community
A	<u>A</u> ccess	Enhance program access for caregivers through timing, location, transportation and publicizing new services
L	<u>L</u> ocal leadership	Include local leadership (nurses, social workers, physicians, ministers, church groups, civic leaders, other community service workers) in supporting and publicizing the program

Using an ecological model of adaptation and aging to view competencies within an environmental context, Lee (1993, pp. 225-227) proposed a number of individual and community-based nursing care approaches. For purposes of this paper they have been modified by the authors to be more germane to service providers in the rural Aging Network, and are set forth in *Exhibit 3*.

Exhibit 3: Individuals and Community-Based Strategies for Caregiver Programs

Individual Strategies	Community-Based Strategies
Because of the heterogeneity of rural environments and the homogeneity of the many subcultures of elderly persons living within those environments, service providers should develop a working knowledge of the characteristics of the local rural environment, the competencies of the elderly persons living there, and the available health and social resources.	Consolidate programs in multi-purpose packages that can serve the broadest possible population. For example, mobile units designed for a specific purpose, such as immunizations, could be expanded to also provide assessment and referral services for elders and their caregivers.
Beyond information on specific physical and mental health conditions, assessment strategies should include information about functional status, individual characteristics, members of kin and non-kin support network, the community environment, and access to transportation. Preferred methods for obtaining information should also be assessed.	Use of existing structures, services or providers as the foundation for developing programs and services for rural caregivers will enhance the acceptability of the programs and increase the longevity of the services.
A home visit should be part of the assessment process, to alert providers to the potential of substandard housing and safety standards that need to be addressed (e.g., a new home safety assessment scale developed by Dr. Louise Poulin de Courval, McGill University: www.clscote-des-neiges.qc.ca/sas).	Linkages between programs need to be formed so that coordination instead of duplication exists at the local level.
Health maintenance and promotion goals should be developed WITH instead of FOR rural elders and their caregivers. Ask what strategies work for them.	Primary care providers (i.e., physicians, nurses, physician's assistants should always be included in the service plan.
Encourage caregivers to use local resources (such as homemaker services) to reduce fatigue secondary to the demands of the caregiving situation. Because asking for and receiving help is not the norm in many rural areas, assist the elder and their caregiver to problem solve while maintaining autonomy, dignity, and privacy. This may require both time and diplomacy.	
Use of control-enhancing interventions may benefit older rural adults and their caregivers. Classes aimed at providing knowledge and skill development (i.e., stress management) may assist caregivers to more adequately master the environment and improve quality of life.	

Betty Grandquist, current Coordinator of the Family Caregiver Support Program in Iowa and former director of the Iowa Department of Elder Affairs, has shared recommendations from her experiences in the aging network, and with rural caregiver support programs in particular. These are:

- ◆ Include volunteers in your program development and implementation efforts. Bring them to the table, as they challenge professionals and offer a fresh and realistic perspective.
- Be sensitive to how the caregiver wants to be perceived...some may reject the "caregiver" label, even though it makes them eligible for services, as they believe being categorized as "caregiver-care recipient" adversely changes the spouse-spouse or adult child-parent relationship.
- ♦ Use a variety of approaches to alert rural caregivers to available services, including brochures, radio and TV spots, educational programs, Web sites, etc. A multi-media consumer publicity "blitz" is a good way to kick off new programs or projects.
- ♦ Many factors are involved in the success or failure of a program in rural settings, such as availability of quality services and transportation.
- Ms. Grandquist advises that flexibility and a common sense approach to the allocation of funds are the hallmarks of any successful program for rural caregivers. For example, in one case a care recipient received burns from an old gas stove while the caregiver was bathing. In this circumstance, an appropriate intervention might be one related to adjustment of the home environment; that is, to use available money to assist the caregiver in purchasing a safer appliance and teaching them how to disable the stove when not directly monitoring the care recipient.
- ◆ To the extent possible, avoid bureaucracies that impede getting the money to where it is needed most.

Dementia-specific recommendations from rural family caregivers in the COEP (Connell et al., 1996, p. 24) included the need to target educational interventions to information and referral agencies (e.g. staff of AAAs), government agencies (e.g. Commission on Aging, law enforcement agencies), and the public using outreach to service clubs, churches, the local business community and community centers. These caregivers also suggested development of a speaker's bureau as a mechanism to disseminate information to the community. Community involvement, a sense of ownership and group identity, and a commitment to the program were viewed as vital to long-term change in the response of people to caregivers of persons with dementia in rural communities.

Finally, Parker et al., (1992, p. 57) offer a number of recommendations on ways to develop case management approaches in rural areas. Their suggestions are based on survey results as well as input from rural case managers in Minnesota and Wisconsin, and were designed to reduce the isolation of rural providers and to increase their support and contact with physicians. Recommendations included: 1) Provide outreach education programs to

professionals; 2) Use the electronic media to disseminate information; 3) Contact congressional officials to reinforce the need to expand Medicare benefits to rural areas; and 4) Improve interdisciplinary teamwork and relationships.

Conclusions

Rural areas have long been characterized as having a unique set of conditions that make service delivery difficult. These include poverty, isolation, difficulties with transportation, sparse and scattered population, resistance to innovation and too few human service agencies, trained professionals and health care resources. Thus, rural health and human service providers are challenged to "define and creatively meet the service needs" (Bice, 1987, p. 9) of rural caregivers. In order to do this, members of the aging network may have to become what Dobkin (2001) has called "social entrepreneurs." That is, persons who "adopt a passion or mission to solve a societal problem" with "continuous creativity, flexibility, and stamina" undeterred by limited resources (Dobkin, 2001, p. 33). To be successful, service providers must offer rural caregivers better coordination of services, improved communication among local agencies (Connell et al., 1996), consistent relationships with providers they trust, and improved access to information (Davis & Magilvy, 2000). Indeed, in order to overcome the many attitudinal and logistic barriers to service delivery in rural areas, community based outreach efforts must include representation from health care professionals, service providers, staff of community organizations and volunteers. "Cooperative efforts help to develop community competence and empowerment, and provide a greater understanding of cultural values and beliefs" (Connell et al., 1996, p. 16).

Effective rural caregiver support programs are those that are caregiver, not provider, driven, and where flexibility is the watchword. Desired programs are available, accessible, accommodating, acceptable and affordable. Some effective programmatic approaches identified in this chapter include: mobile outreach programs, in-home visitation, satellite clinics, and increased use of telemedicine and information technologies. The RURAL model is set forth as a way to provide caregiver support and training and to meet the diverse needs of rural caregivers while accommodating the characteristics of their social networks.

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RECOMMENDED RESOURCES

1. For information on elder care resources, including specific information for caregivers, see:

www.resinets.com/health/elder.htm

2. For information on general health issues in rural America, contact:

www.nrharural.org

3. For activities of all state offices of rural health, contact:

www.rural.center.org/nosorh/

4. For further information about rural case management, see:

Parker, M., Quinn, J., Viehl, M., McKinley, A.H., Polich, C.L., Hartwell, S., Van Hook, R., and Detzner, D.F. (1992). Issues in Rural Case Management. *Family and Community Health*, 14(4), 40-60.

5. For Information Regarding Assessment Issues for Rural Case Management, and How to Blend Extant AAA Data Bases contact:

Jon Lemke, PhD University of Iowa Center on Aging 2159 Westlawn

Iowa City, IA 52242 Ph#: (319) 335-7569

E-Mail: jon-lemke@uiowa.edu

6. For Information on Faith-Based Communities, the Volunteer Caregiver Program at the University of Louisville, and Social Skills Training Groups for Caregivers contact:

Karen Robinson, RN, PhD, FAAN

University of Louisville School of Nursing

555 South Floyd Street

Louisville, KY 40292

Ph#: (502) 852-8512

E-Mail: kmrobi01@louisville.edu

7. For Information Regarding the New Home Safety Assessment Scale for People with Dementia Living at Home contact:

Louise Poulin de Courval

McGill University Ph#: (514) 731-1386

E-Mail: mclp@musica.mcgill.ca, or see www.clsccote-des-neiges.qc.ca/sas/

Administration on Aging ADDGS Rural Iowa Demonstration Project 2000-2001 Grant number No. 90AZ2366 Second Edition

A1. The Client Family	ID# <u>2 0</u>				
A2. Date of intake	/				
	Month	day	year		
A3. County ID#	Boone	08			
(circle response)	Cedar	16			
	Clinton	23			
	Ida	47			
	Iowa	48			
	Madison	61			
	Monona	67			
	Muscatine	70			
A4. Interviewer's initi	als				
	THE CLL	ENT INT	TAKE FORM	1	
NOTE E 4: 0					1.1
				al with memory difficults along the for The Client's ca	
CAREGIV	EK, refers to the p	erson mo	si responsio	ite for the Chefit's ca	116.
THE CLIENT NAME					
THE CLIENT NAME					
	First name		Surname		
ADDRESS					
City	State				
ZIP CODE	PHO	NE()		
Zip (Require	d)				
CAREGIVER NAME					
(Name of person most responsible for	or the Client's care) First t			Surname	
	or the cheft search That I	141110		Samuno	
ADDRESS	Q. :				
City	State				
ZIP CODE	PHONE () _		<u> </u>	

PART 1: INFORMATION ABOUT THE CLIENT

Client Demographics	Form D		
Instructions: Circle response			
D1 . THE CLIENT gender: Male Female 1			
D2. THE CLIENT's marital status: single married/domestic partner widowed other 1 married/domestic partner 2 widowed 3 other 4			
D3. THE CLIENT's birth date:			
D4. Does THE CLIENT have Alzheimer's disease? Yes, (Alzheimer's disease or other dementia has been medically diagnosed Probably, (Alzheimer's disease or other dementia is suspected) No (Record other diagnosis if known)) 1 2 3		
D5. How do you know? suspected not diagnosed diagnosis by family MD diagnosis by specialist diagnosis by specialized clinic other (specify) suspected not diagnosed 1 diagnosed 2 diagnosis by specialized 5			
D6. Geographic location of <i>THE CLIENT's</i> residence: a rural or farm community (fewer than 2,500 people) small city or town that is not suburb of a larger city (2,500 to 50,000 people a medium sized city or suburb of medium size city (50,000 to 100,000 people a large city or suburb of large city (more than 100,00 people) an Indian reservation	*		
D7. Where does THE CLIENT reside? Lives in house or apartment with others(s) (go to 8) Lives alone in house or apartment (skip to 9) Lives in a group environment with assistance (skip to 9) (not a nursing home) Lives in nursing home (skip to 9) Other	1 2 3 4 5		
D8. If THE CLIENT lives in house or apartment, how many people reside in the household? (Include THE CLIENT in the total number.) Total Persons in household			

D9. How much help, if any, does THE CLIENT need with each of these activities?

	needs no help/no supervision	need some help/occasional supervision	needs a lot of help/constant supervision	can't do it at all
(a) Eating	1	2	3	4
(b) Getting in and out of bed	1	2	3	4
(c) Getting around inside	1	2	3	4
(d) Dressing	1	2	3	4
(e) Bathing	1	2	3	4
(f) Using the toilet	1	2	3	4
(g) Doing heavy housework	1	2	3	4
(h) Doing light housework	1	2	3	4
(i) Doing laundry	1	2	3	4
(j) Cooking/preparing meals	1	2	3	4
(k) Buying/getting food/clothes	1	2	3	4
(l) Getting around outside	1	2	3	4
(m) Going places outside of walking distance	1	2	3	4
(n) Managing money	1	2	3	4
(o) Taking medicine	1	2	3	4
(p) Using telephone	1	2	3	4
(q) Driving	1	2	3	4

D10. How did you learn of this program?

1 0	
mass media	1
physician	2
brochure	3
friend	4
church religious organization	5
service agency or case manager	6

D11. Name or person/ agency/ organization

D12. Which of these categories is closest to *THE CLIENT's* total annual income? (If THE *CLIENT* is married include income of spouse)

under \$8,000	1
\$8,000 - \$11,999	2
\$12,000 - \$14,999	3
\$15,000 - \$19,999	4
\$20,000 - \$29,999	5
\$30,000 and over	6

D13. Is the Elder of Spanish, Hispanic or I	Latino origin?	
No Yes (Record group, e.g. Mexican, Chio	cano, Cuban)	1 2
Too (Iteoora group, e.g. Irromoun, e.m.		<i>_</i>
D14. Which of the following categories be	st describes Elder's race?	
White Black, African-American or Negro American Indian or Alaska Native (Re Asian (Record Race) Pacific Islander (Record Race)		1 2 3 4 5
D15. In which language(s) is the CLIENT fl (check all that apply)	luent? English Spanish Other, (list)	1 2 3
D16. Does the CLIENT still drive?	Yes	1
	No	2
D17. Does the CLIENT family have a car?	Yes	1
	No	2

Health status - The Client

Form HP

Instructions; *To be asked of the Client. Circle the response given.*

HP1. Compared to other people your age, would you say that your general health is?

Excellent	1
Good	2
Fair	3
Poor	4
Very poor	5
Don't know	6

HP2. Has your general health changed much in the last 12 months?

Much better	1
Somewhat better	2
About the same	3
Somewhat worse	4
Much worse	5
Don't know	6

HP3a) Has the Client seen a doctor or other health care practitioner since we last spoke?

Y / N

b) How many times?

1 2 3 4 5 6 >6

HP4a) Has the Client been in hospitalization at least overnight since we last spoke?

Y / N

b) How many times?

1 2 3 4 5 6 >6

c) For what length of stay (#days) ____ ___

HP5a) Has the Client been in placed in a nursing home at least overnight since we last spoke?

Y / N

b) How many times?

1 2 3 4 5 6 >6

c) For what length of stay (#days) ____ __

ADL/ IADL functional abilities

Form AD

Instructions; Each of the following questions is in two parts. Please rate each on the following 1-5 scale, then answer each Y or N;

1 = just unable to do it 2= a lot of difficulty 3= some difficulty 4= a little difficulty 5 = no difficulty 9= don't know

- AD1. How <u>difficult</u> on average is it for the Client to do each of the following kinds of activities without any assistive device?
- AD2. Is the activity possible with an assistive device of some sort?

				A		Don't		
<u>Difficulty</u>	Unable	A lot	Some	little	None	know	Dev	ice?
a. lifting or carrying groceries	1	2	3	4	5	9	Y	N
b. reading comprehension	1	2	3	4	5	9	Y	N
c. climbing one flight of stair	1	2	3	4	5	9	Y	N
d. bending, kneeling, or stooping	1	2	3	4	5	9	Y	N
e. writing	1	2	3	4	5	9	Y	N
f. bathing	1	2	3	4	5	9	Y	N
g. grooming	1	2	3	4	5	9	Y	N
h. selecting clothing	1	2	3	4	5	9	Y	N
i. dressing yourself	1	2	3	4	5	9	Y	N
j. handling or fingering small objects (buttons, zippers, eating utensils)	1	2	3	4	5	9	Y	N
k. using the toilet	1	2	3	4	5	9	Y	N
l. walking more than a mile (outdoors)	1	2	3	4	5	9	Y	N
m. walking several blocks (outdoors)	1	2	3	4	5	9	Y	N
n. walking around residence	1	2	3	4	5	9	Y	N

Global Deterioration Scale

Form GD

Date	/	/	1		

Instructions: From the table below select the <u>one</u> GDS Stage that best fits this care recipient. Circle the number for that stage on the GDS Score line below.

GD1. GDS Score 1 2 3 4 5 6 7

GDS STAGE	CLINICAL PHASE CLINICAL CHARACTERISTICS
1 = NO COGNITIVE DECLINE (Normal)	 No subjective complaints of memory deficit. No memory deficit evident on clinical interview.
2 = VERY MILD COGNITIVE DECLINE (Forgetfulness)	 Subjective complaints of memory deficit, most frequently in following areas: a. forgetting where one has placed familiar objects; and, b. forgetting names one formerly knew well. No objective deficits in employment or social situations. Appropriate concern with respect to symptomatology.
3 = MILD COGNITIVE DECLINE (Early Confusional)	 Earliest clear-cut deficits. Manifestations in more than one of the following areas: a. a patient may have gotten lost while traveling to an unfamiliar location; b. co-workers become aware of patient's relatively poor performance;
	 c. word and name finding deficits become evident to intimates; d. patient may read a passage or a book and retain relatively little material; e. patient may demonstrate decreased facility in remembering names upon introduction to new people; f. patient may have lost or misplaced an object of value; g. concentration deficit may be evident on clinical testing. Decreased performance in demanding employment and social setting. Denial begins to become manifest in patient. Mild to moderate anxiety accompanies symptoms.
4 = MODERATE COGNITIVE DECLINE (Late Confusional)	 Clear-cut deficit on careful clinical interview. Deficits manifest in following areas: decreased knowledge of current world events and recent events in own life; may exhibit some deficit in memory of personal history; concentration deficit elicited on serial subtractions; decreased ability to travel, handle finances, etc. Frequently no deficit in the following areas: orientation to time and person; recognition of familiar persons and faces; ability to travel to familiar locations. Inability to perform complex tasks. Denial is dominant defense mechanism. Flattening of affect and withdrawal from challenging situations occur.

5 = MODERATELY SEVERE COGNITIVE DECLINE (Early Dementia)

- Patients can no longer survive without some assistance.
- Patients are unable during interview to recall a major relevant aspect of their current lives, e.g.:
 - a. their address or telephone number of many years;
 - b. the names of close members of their family (such as grandchildren);
 - c. the name of the high school or college from which they graduated.
- Frequently some disorientation to time (date, day of week, season, etc.) or to place.
- An educated person may have difficulty counting back from 40 by 4s or from 20 by 2s.
- Persons at this stage retain knowledge of many major facts regarding themselves and others. They invariably know their own names and generally know their spouse's and children's names.
- They require no assistance with toileting or eating, but may have some difficulty choosing the proper clothing to wear.

6 = SEVERE COGNITIVE DECLINE (Middle Dementia)

- May occasionally forget the name of the spouse whom they are entirely dependent upon for survival. Will be largely unaware of all recent events and experiences in their lives.
- Retains some knowledge of their past lives but this is very sketchy. Generally unaware of their surroundings, the year, the season, etc.
- Almost always recalls own name. Frequently continue to be able to distinguish familiar from unfamiliar persons in their environment.
- May have some difficulty counting from 10, both backward and forward.
- Will require some assistance with activities of daily living, e.g., may become incontinent. Will require travel assistance but occasionally will display ability to travel to familiar locations. Diurnal rhythm frequently disturbed.
- Personality and emotional changes occur. These are quite variable and include:
 - a. delusional behavior, e.g., patients may accuse their spouse of being an imposter; may talk to imaginary figures in the environment, or to their own reflection in the mirror;
 - b. obsessive symptoms, e.g., persons may continually repeat simple cleaning activities;
 - c. anxiety symptoms, agitation, and even previously nonexistent violent behavior may occur;
 - d. cognitive abulia, i.e., loss of will power because an individual cannot carry a thought long enough to determine a purposeful course of action.

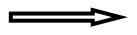
7 = VERY SEVERE COGNITIVE DECLINE (Late Dementia)

- All verbal abilities are lost. Frequently there is no speech at all only grunting.
- Incontinent of urine; requires assistance toileting and feeding.
- Loses basic psychomotor skills, e.g., ability to walk. The brain appears no longer able to tell the body what to do. Generalized and cortical neurologic signs and symptoms are frequently present.

ans	Date_ce a check (\$\) for each correct answer. Leave wers blank. Score one point for each blank. Abtotal) and subtract from 30 for the score.	
1. What is the year_	, season, date, month	1. ()
2. Where are we: sta	ate, county, town, street, number	2. ()
3. Name 3 objects: o	orange, airplane, tobacco (trials).	3. ()
4. Serial 7's(9 OR spell "world"	73),(86),(79),(72),(65). backwards(d)(l)(r)(o)(w).	4. ()
5. Recall 3 objects:	orange, airplane, tobacco	5. ()
	, and watch	6. ()
(Show the objects 7. Read and obey	S). CLOSE YOUR EYES	7. ()
(if they have poor 8. Copy design	r eyesight, this question is enlarged below) (below).	8. ()
9. Write a sentence	(below).	9. ()
10. Repeat the follo	wing "no ifs, ands, or buts"	10.()
11. Follow a 3 stage	b. fold it in half c. put it on the floor	11.()
		SUBTOTAL
Design to copy:		
MM1.Level of conso	ciousness: Alert=1 Drowsy =2 Stupor=3 Coma=4	I

MM2MMSE Score
(30- subtotal= score)

7. Read and obey.



CLOSE YOUR EYES

Behavior Rating Checklist

Form BR

Date	/	/	
------	---	---	--

Instructions; Rate and circle each item on a scale of 1 to 5, I = extremely or daily, 2 = much or weekly, 3 = moderately, or 2-3 times per month, 4 = slightly or monthly, 5 = absent, or 9 = don't know.

In the last month, have these symptoms or behaviors occurred;

						Don't
	Extremely	Much	Moderately	Slightly	Absent	know
1. Anxiety	1	2	3	4	5	9
2. Restlessness	1	2	3	4	5	9
3. Tense	1	2	3	4	5	9
4. Seems afraid of something	1	2	3	4	5	9
5. Becomes easily upset	1	2	3	4	5	9
6. Seeks reassurance from others	1	2	3	4	5	9
7. Hostility	1	2	3	4	5	9
8. Unfriendly to others	1	2	3	4	5	9
9. Impolite to others	1	2	3	4	5	9
10. Complains	1	2	3	4	5	9
11. Objects to some routine procedures	1	2	3	4	5	9
12. Angry	1	2	3	4	5	9
13. Depressed	1	2	3	4	5	9
14. Seems to feel rejected	1	2	3	4	5	9
15. Sad appearance	1	2	3	4	5	9
16. Withdrawn	1	2	3	4	5	9
17. Quiet	1	2	3	4	5	9
18. Talks of gloomy things	1	2	3	4	5	9
19. Calm	1	2	3	4	5	9
20. Cheerful	1	2	3	4	5	9
21. Friendly	1	2	3	4	5	9
22. Seems confused	1	2	3	4	5	9
23. Increased confusion at night late day /	1	2	3	4	5	9
24. Lacks recognition of significant others	1	2	3	4	5	9
25. Aggressiveness	1	2	3	4	5	9
26. Hallucinations	1	2	3	4	5	9
27. Paranoid thoughts	1	2	3	4	5	9
28. Yells inappropriately	1	2	3	4	5	9
29. Sexual acting out	1	2	3	4	5	9
30. Wanders away	1	2	3	4	5	9

Functional Assessment II

Form FA

Categories A - D

- Categories A D are considered override categories.
- A level three rating in any one-override category indicates a Level 3 designation regardless of a care recipient's total score on the remainder of the categories.

All Criteria Have Equal Value

- Many of the descriptors under each heading (listed with serial numbers) have more than one criterion.
- All criteria have equal value. The time criterion should not take precedence over the other listed criteria.
- A care recipient does not have to meet all the criteria listed under the corresponding number to be considered functioning at that level.
- A care recipient's score (rating) in each area should capture the overall type of assistance needed by the care recipient.

All Categories Must Be Scored

• Circle a score in all categories.

Instrument Constraints

- The Functional Assessment II is not intended to replace clinical or professional assessment.
- The FA II is designed as a tool that captures **caregiver** time and/or skill required to meet care recipient care needs.

Instructions: Rater needs to have direct observation of care recipient prior to completing level designation. Circle rating interval and level 1,2, 3 for each indicator.

Visit 1) INITIAL 2) 6 MO 3) 12 MO 4) 18 MO 5) 24 MO

CURRENT DATE	/	/		
A. HEALTH RELATED CONDITION	1	2	3	
B. CARE MANAGEMENT	1	2	3	
C. COMMUNICATION	1	2	3	
D. ANXIETY	1	2	3	
E. Dietary	1	2	3	
F. Eating (Facilitation Of)	1	2	3	
G. Restorative Therapies	1	2	3	
H. Toileting	1	2	3	
I. Ambulating / Mobility	1	2	3	
J. Social Interaction	1	2	3	
K. Medications	1	2	3	
L. Hygiene	1	2	3	
TOTAL (max. 12)				
LEVEL	1	2	3	

FUNCTIONAL ASSESSMENT II INDICATORS

A. HEALTH RELATED CONDITION:

(Safety, medical conditions, mental health, and cognitive deficits).

- 1. Health conditions multiple but stable. Baseline routine maintained with routine screening and assessment.
- 2. Health condition requires ongoing planned or intermittent monitoring and intervention. Required once per week.
- 3. Fluctuating/variable or complex condition(s) require regular monitoring or intervention of a health professional. Requires monitoring more than once a week.

B. CARE MANAGEMENT:

(Coordinating activities on behalf of the care recipient either with family, care partners or other agencies or care providers).

- 1. Care recipient, family able to coordinate and address care needs and issues. Less than 1 https://month.nih.gov/html in contact with health care professional or other caregivers regarding care recipient condition/concerns. Regular scheduled appointments captures majority of needs.
- 2. 2 <u>hr./month contact</u> with health care professional or other caregivers regarding care recipient condition/concerns <u>or</u> < 2 <u>unplanned</u> contacts per month to care provider or other health care professional:
 - assistance with managing behavioral limitations (e.g. accommodating the environment)
 - assistance with implementation of health regimes
 - assistance in coordinating ancillary medical/social and support services
 - assistance in coordinating financial arrangements/ assistance (e.g., VA and Elderly Wavier)
- 3. 2+ <u>hr./month</u> contact with or on behalf of family/caregiver regarding care recipient condition/concerns or > 2 <u>unplanned</u> contacts per month or continuous caregiver demands 1+ hours per month:
 - Recurring issues and/or non-compliance with agreed intervention
 - day center not kept informed of issues effecting care here.

C. COMMUNICATION:

(Impaired reception or expression regarding needs, feelings, frustration. Result from neuro, psych, cognitive, sensory deficit).

- 1. Verbalizes or gestures to express needs so as to be understood in most situations.
- 2. Needs caregiver intervention to clarify basic needs and ideas ≤ 15 minutes per day.
- 3. Needs caregiver intervention to clarify basic needs and ideas 15+ minutes per day:
 - unable to express needs appropriately or consistently despite prompting.

D. ANXIETY:

(A condition of mental uneasiness arising from fear, solicitude or apprehension)

- 1. Requires minimal reassurance and/or redirection;
 - responds to planned routine
 - engages with others
 - passively occupied with environment props in place (towels, puzzles, cards)
- 2. Responds to moderate reassurance and/or redirection;
 - able to be engaged to prevent crisis by planned interventions
 - 0 15 minutes per day.
- 3. Requires continuous reassurance and/or redirection;
 - unplanned interventions
 - unpredictable elopement
 - unresponsive to redirection and retrieval
 - 15+ minutes per day.

E. DIETARY:

- 1. Nutrition maintained by food delivery or home meal preparation.
- 2. Routine substitutions and supplements, follows dietary guideline without intervention.
- 3. Special preparation and alteration by caregiver, unplanned interventions (supplements, walking foods, policing and cueing for dietary compliance).

F. EATING: (Facilitation Of)

- 1. Eats independently, no risk for choking or aspiration.
- 2. Assistance with cutting, special utensils, setting adaptation.
- 3. Constancy of monitoring or companion to assure nutritional intake/safety, or strategic placement to facilitate eating or minimize disruption.

G. RESTORATIVE THERAPIES:

- 1. Routine followed by self-initiation.
- 2. Routine requires prompting to initiate.
 Individual's routine requires monitoring to integrate into daily routine.
 Requires one-on-one, ≤ 15 minutes per day.
- 3. Requires caregiver direction may be one-on-one; ≥15 minutes per day.

H. TOILETING:

- 1. No continence issues, or tends to needs with reminders.
- 2. Continent with bowel and bladder program, ≤ 15 minutes per day (time may include perennial hygiene and hand washing).
- 3. Incontinent (unplanned), 15+ min. per day.

I. AMBULATION / MOBILITY:

- 1. Ambulates/transfers independently with or without device, follows safety precautions.
- 2. Mobility and transfers managed at transition times with stand-by supervision or hands on assistance. Follows safety precautions with cueing, ≤15 minutes.
- 3. Requires assistance of one or more people, unplanned frequent monitoring or intervention, 15+ minutes.

J. SOCIAL INTERACTION:

(Individual's input coincides with group conversation and/or activity).

- 1. Interacts appropriately with **caregiver**, **other family**, **and friends** (e.g., controls temper, is aware that words and actions affect others, makes appropriate requests).
- 2. Requires **caregiver** intervention ≤ 15 minutes/day:
 - participates in or on the edge of the group
 - has trouble in unusual situations
 - makes frequent requests. Or lack of requests
 - little if any benefit from others.
- 3. Requires **caregiver** intervention 15+ minutes/day:
 - requires supervision, e.g., monitoring, cueing, or coaxing.

K. MEDICATIONS:

- 1. Takes no meds or takes meds independently with cueing.
- 2. Medications set-up by home caregiver:
 - requires oversight for administration x 2 daily
 - medications stable
 - stand-by supervision, cueing, or coaxing ≤ once/day.
- 3. 24 hour set-up requires professional assistance to caregiver:
 - requires oversight for administration x 3 daily
 - medication changes frequently
 - intermittent or PRN medications
 - stand-by supervision, cueing, or coaxing more than once a day.

L. HYGIENE and GROOMING:

(hands, ADL, infection risk, shower)

- 1. Meets own hygiene needs. Grooming is non-offensive. Occasional verbal prompt.
- 2. Manages with schedule, occasional reminders:
 - nail and hair care, soiled clothing, requires caregiver ≤ 15 minutes per day
 - planned shower or bath
 - indiscriminate touching (infection control practices manage)
 - responds to verbal cues for nose blowing, coughing, hand washing.
- 3. Requires assistance with hygiene 15 + minutes per day:
 - indiscriminate touching of objects requires monitoring 15+ per day.

PART 2: INFORMATION ABOUT CAREGIVER

(Person most responsible for The Client's care)

Caregiver Demographics/Status

Form CG

Note: Do **not** record this information if the caregiver is a professional and unrelated to the client. Instructions: Circle response **CG1**. *CAREGIVER'S* relationship to the *CLIENT*: 1 no caregiver identified spouse/domestic partner 2 child/child-in-law 3 sibling 4 other relative 5 friend/neighbor 6 professional care manager 7 CG2. How long has CAREGIVER provided most of the care? less than 6 months 1 6 to 12 months 2 to 24 months 3 25 months to 5 years 4 more than 5 years 5 CG3. CAREGIVER'S birth date? Month Day Year CG4. CAREGIVER'S gender? Male 1 Female 2 CG5. CAREGIVER'S marital status? single 1 married/domestic partner 2 widowed 3 other 4 CG6. Is the CAREGIVER of Spanish, Hispanic or Latino origin? No 1 Yes 2 (Record group, e.g. Mexican, Chicano, Cuban) CG7. Which of the following categories best describes Client's race? White Black, African-American or Negro American Indian or Alaskan Native (Record principal tribe) 3 Asian (Record Race) 4 Pacific Islander (Record Race)

CG8. In which language(s) is the CAREGA	IVER fluent?	English	1
(check all that apply)		Spanish	2
		Other,	3
		(list),	
CG9. CAREGIVER's employment status:	works full-tir	ne	1
1 0	works part-tii		2
		orks part-time	3
	fully retired	1	4
	homemaker		5
	unemployed		6
	other		7
		THE I A	,
CG10. What is the highest grade in school	ol that CAREG	IVER complete	
8 th grade or less			1
attended high school	(D: 1	IED)	2
high school graduate			3
some college or post		uning	4
Associate degree (AA			5
Bachelor's degree (B	S, BA, etc.)		6
graduate degree			7
CG11. During the past week, about how CLIENT with (if not a previous res	•	total did the	CAREGIVER help THE
11a. Eating, bathing, dressing or helping	g with toilet fur	nctions	(#hours per week)
11b. Meal preparation, laundry or light h	housework		(#hours per week)
11c. Providing transportation to appoint	ments and/or sl	hopping	(#hours per week)
11d. Legal matters, banking or money n	natters		(#hours per week)
CG12. Approximately how far away in C	0		
CLIENT? (0 minutes, if caregiver		-	 ·
CG13. Which of the following services is	the CLIENT F	AMILY curren	tly using?
			(ADEGMED)

(Circle $\underline{\mathit{all}}$ services that are used by either the CLIENT OR the $\mathit{CAREGIVER}$)

a.	companion or friendly visitor	Yes	No
b.	transportation services	Yes	No
c.	supervision	Yes	No
d.	case management	Yes	No
e.	homemaker services	Yes	No
f.	support groups	Yes	No
g.	chore services	Yes	No
h.	caregiver training program	Yes	No
i.	personal care services	Yes	No
j.	psychological counseling	Yes	No
k.	home health services	Yes	No
1.	group meals/home delivered meals	Yes	No
m.	adult daycare center/ adult day health	Yes	No
n.	respite in a nursing home, adult foster	Yes	No
	home, or someone else's home		
0.	Other service, list	Yes	No

CG14. Client support- What do you see that you need help with?

a)	Personal care	Yes	No
b)	Transportation	Yes	No
c)	Personal finance	Yes	No
d)	Adult day care	Yes	No
e)	Congregate meals	Yes	No
f)	Social contacts	Yes	No
g)	Housekeeping	Yes	No
h)	Shopping/ errands	Yes	No
i)	Services from a health professional	Yes	No
j)	Home delivered meals	Yes	No
k)	Emergency response system	Yes	No
1)	Telehealth visits	Yes	No

CG15. Which of these categories is closest to the CAREGIVER's total annual HOUSEHOLD income excluding any income of the CLIENT?

under \$8,000	1
\$8,000 - \$11,999	2
\$12,000 - \$14,999	3
\$15,000 - \$19,999	4
\$20,000 - \$29,999	5
\$30,000 - \$39,999	6
\$40,000 and over	7

CG16. Does THE CAREGIVER drive?

Yes	1
No	2

Health status - Caregiver

Form HC

Instructions; To be asked of the caregiver. Circle the response given.

HC1. Compared to other people your age, would you say that your general health is?

Excellent	•••••	1
Good	•••••	2
Fair	•••••	3
Poor	•••••	4
Very poor	•••••	5
Don't know	•••••	6

HC2. Has your general health changed much in the last 12 months?

Much better	•••••	1
Somewhat better	•••••	2
About the same	•••••	3
Somewhat worse	•••••	4
Much worse		5
Don't know	•••••	6

HC3a) Has the Caregiver seen a doctor or other health care practitioner since we last spoke?

b) How many times?

HC4a) Has the Caregiver been in hospitalization at least overnight since we last spoke?

- b) How many times?
- 1 2 3 4 5 6 >6
- c) For what length of stay (#days) ____

HP5a) Has the Caregiver been in placed in a nursing home at least overnight since we last spoke?

- b) How many times?
- 1 2 3 4 5 6 >6
- c) For what length of stay (#days) _____

Form S

<u>Definition:</u> The extent of bio-psycho-social pressure on a family care provider caring for a family member or significant other over an extended period of time.

Instructions: Rate and circle each indicator on a scale of 1 to 5; 1 = Extensive to 5 = None. Then give an overall rating using the same scale.

Can	ogivon Stuossons	Extensive	Substantial	Moderate	Limited	None
	egiver Stressors					
S1	Reported stressors of caregiving	1	2	3	4	5
S2	Physical limitations for caregiving	1	2	3	4	5
S3	Psychological limitations for caregiving	1	2	3	4	5
54	Cognitive limitations for caregiving	1	2	3	4	5
S5	Impairment of usual role performance	1	2	3	4	5
36	Impairment of social interactions	1	2	3	4	5
S7	Perceived lack of social support	1	2	3	4	5
38	Perceived lack of health care system support	1	2	3	4	5
S9	Lack of usual diversional activity	1	2	3	4	5
S10	Impairment of usual work performance	1	2	3	4	5
S11	Severity of care recipient illness	1	2	3	4	5
S12	Amount of care or oversight required	1	2	3	4	5
S13	Impairment of caregiver- patient relationship	1	2	3	4	5
514	Other(Specify)	1	2	3	4	5
S15	Overall rating	1	2	3	4	5

Caregiver Well-Being

Form W

<u>Definition</u>: Primary care provider's satisfaction with health and life circumstances *Instructions*: Rate and circle each indicator on a scale of 1 to 5; 1 = Extremely compromised to 5 = Not compromised.

Then give an overall rating using the same scale.

		Extremely compromised	Substantially compromised	Moderately compromised	Mildly compromised	No compromise
Careg	giver Well-Being					
W1.	Satisfaction with physical health	1	2	3	4	5
W2.	Satisfaction with emotional health	1	2	3	4	5
W3.	Satisfaction with lifestyle	1	2	3	4	5
W4.	Satisfaction with performance of usual roles	1	2	3	4	5
W5.	Satisfaction with social support	1	2	3	4	5
W6.	Satisfaction with instrumental support	1	2	3	4	5
W7.	Satisfaction with professional support	1	2	3	4	5
W8.	Satisfaction with social relationships	1	2	3	4	5
W9.	Satisfaction with caregiver role	1	2	3	4	5
W10	Other	1	2	3	4	5
	(Specify)					
W11	Overall rating	1	2	3	4	5

Caregiving Endurance Potential

Form E

<u>Definition</u>: Factors that promote family care provider continuance over an extended period of time

Instructions: Rate and circle each indicator on a scale of 1 to 5; 1 = Not adequate to 5 = Totally adequate. Then give an overall rating using the same scale.

		Not adequate	Slightly adequate	Moderately adequate	Substantially adequate	Totally adequate
Caregi	iving Endurance Potential					
E1.	Mutually satisfying care recipient-caregiver relationship	1	2	3	4	5
E2.	Mastery of direct care activities	1	2	3	4	5
E3.	Mastery of indirect care activities	1	2	3	4	5
E4.	Services needed for the care recipient	1	2	3	4	5
E5.	Social support for caregiver	1	2	3	4	5
E6.	Health care system support for caregiver	1	2	3	4	5
E7.	Resources to provide care	1	2	3	4	5
E8.	Respite for caregiver	1	2	3	4	5
E9.	Opportunities for caregiver leisure activities	1	2	3	4	5
E10.	Other(Specify)	1	2	3	4	5
E11.	Overall rating	1	2	3	4	5